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Malcolm Wilson MOON et al.

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FOREIGN PATENT DOCUMENTS

	REF	DOCUMENT NUMBER	DATE	COUNTRY	CLASS	SUB- CLASS	TRANSLATION	
							YES	NO
RA	A1	99/61422	12/02/1999	WO	—	—		
RA	A2	00/08202	02/17/2000	WO	—	—		
RA	A3	01/90104	11/29/2001	WO	—	—		

OTHER DOCUMENTS (Including Author, Title, Date, Pertinent Pages, Etc.)

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Form PTO-1449 (MODIFIED)	U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE	ATTY. DOCKET NO. 038602/1153	SERIAL NO. 09/863,804
INFORMATION DISCLOSURE CITATION (Use several sheets if necessary)		APPLICANT MOON et al.	
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U.S. PATENT DOCUMENTS

EXAMINER INITIAL	REF	DOCUMENT NUMBER	DATE	NAME	CLASS	SUB- CLASS	FILING DATE IF APPROPRIATE
RA	A1	2968557	1/17/61	Burgandt et al.	—	—	
RA	A2	4002749	1/11/77	Rovnyak	—	—	
RA	A3	4053613	10/11/77	Rovnyak et al.	—	—	
RA	A4	4642309	2/10/87	Michel et al.	—	—	
RA	A5	4826847	5/2/89	Michel et al.	—	—	
RA	A6	5051417	9/24/91	Nadler et al.	—	—	
RA	A7	5124347	6/23/92	Connor et al.	—	—	
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RA	A10	5322950	6/21/94	Sircar et al.	—	—	
RA	A11	5374652	12/20/94	Buzzetti et al.	—	—	
RA	A12	5382593	1/17/95	Le Baut et al.	—	—	
RA	A13	5389661	2/14/95	Sircar et al.	—	—	
RA	A14	5397787	3/14/95	Buzzetti et al.	—	—	
RA	A15	5409949	4/25/95	Buzzetti et al.	—	—	
RA	A16	5792783	8/11/98	Tang et al.	—	—	
RA	A17	5834504	11/10/98	Tang et al.	—	—	
RA	A18	5849710	12/15/98	Battistini et al.	—	—	
RA	A19	5880141	3/9/99	Tang et al.	—	—	
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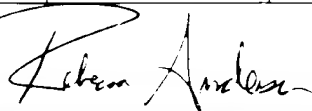
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7A	A22	5886020	3/23/99	Tang et al.	—	—	
2A	A23	6133305	10/17/00	Tang et al.	—	—	

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EXAMINER INITIAL	REF	DOCUMENT NUMBER	DATE	COUNTRY	CLASS	SUB- CLASS	TRANSLATION	
							YES	NO
KA	A24	91/13055	9/5/91	WO	—	—		
RA	A25	92/07830	5/14/92	WO	—	—		
RA	A26	92/20642	11/26/92	WO	—	—		
RA	A27	93/01182	1/21/93	WO	—	—		
RA	A28	94/14808	7/7/94	WO	—	—		
RA	A29	95/01349	1/12/95	WO	—	—		
RA	A30	95/17181	6/29/95	WO	—	—		
RA	A31	96/00226	1/4/96	WO	—	—		
RA	A32	96/16964	6/6/96	WO	—	—		
RA	A33	96/22976	8/1/95	WO	—	—		
RA	A34	96/32380	10/17/96	WO	—	—		
RA	A35	96/40116	12/19/96	WO	—	—		
RA	A36	97/25986	7/24/97	WO	—	—		
RA	A37	98/07695	2/26/98	WO	—	—		
RA	A38	98/24432	6/11/98	WO	—	—		
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RA	A42	99/52869	10/21/99	WO	—	—		
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RA	A44	99/65869	12/23/99	WO	—	—		
RA	A45	00/08202	2/17/00	WO	—	—		
RA	A46	00/35906	6/22/00	WO	—	—		
RA	A47	00/35908	6/22/00	WO	—	—		
RA	A48	00/35909	6/22/00	WO	—	—		
RA	A49	00/38519	7/6/00	WO	—	—		
RA	A50	00/56709	9/28/00	WO	—	—		
RA	A51	01/60814	8/23/01	WO	—	—		
RA	A52	2159360	11/30/71	Germany	—	—		
RA	A53	2159361	11/30/71	Germany	—	—		
RA	A54	2159362	11/30/71	Germany	—	—		
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RA	A56	2321656	4/28/73	Germany	—	—		
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				MOON et al.			
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RA	A69	2689397	10/8/93	France	—	—	
RA	A70	809691	3/4/59	Great Britain	—	—	
RA	A71	835473	5/18/60	Great Britain	—	—	
RA	A72	6229570	7/2/87	Japan	—	—	
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RA	A79	Andreani et al., "Synthesis and cardiotoxic activity of 2-indolinones," <u>Eur. J. Med. Chem.</u> 25:187-190 (1990)					
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RA	A81	Andreani et al., "Synthesis and cardiotoxic activity of pyridylmethylene-2-indolinones," <u>Eur. J. Med. Chem.</u> 27:167-170 (1992) © Elsevier, Paris					
RA	A82	Andreani et al., "Synthesis and potential coanthracyclinic activity of substituted 3-(5-imidazo[2,1-b]thiazolylmethylene)-2-indolinones," <u>Eur. J. Med. Chem.</u> 32:919-924 (1997) © Elsevier, Paris					
RA	A83	Andreani et al., "Synthesis of lactams with potential cardiotoxic activity," <u>Eur. J. Med. Chem.</u> 28:825-829 (1993)					
RA	A84	Andreani et al., "In Vivo Cardiotoxic Activity of Pyridylmethylene-2-indolinones," <u>Arzneimittel-Forschung Drug Research</u> 48:727-729 (1998) ©					
RA	A85	Bahner and Brotherton, "9-(4-Aminobenzylidene)fluorenes," <u>J. Med. Chem.</u> 12:722-723 (1969)					
RA	A86	Bahner et al., "Benzylideneindenes with Oxygen Attached to the Indene Ring," <u>J. Med. Chem.</u> 12:721-722 (1969)					
RA	A87	Bamfield et al., "Diels-Alder Reactions of Oxindolylideneacetone," <u>J. Chem. Soc. (C)</u> 1028-1030 (1966) ©					
RA	A88	Borsche et al., "Über vielkernige kondensierte Systeme mit heterocyclischen Ringen. XIII.," <u>Liebigs Ann. Chem.</u> 550:160-174 (1941)					

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RA	A89	Buzzetti et al., "Cinnamamide Analogs as Inhibitors of Protein Tyrosine Kinases," <u>Il Farmaco</u> 48:615-636 (1993)	
RA	A90	Chatten et al., "Substituted Oxindoles. Part VI. Polarographic Reduction of Substituted <i>trans</i> -3-Benzylideneindol-2(3 <i>H</i>)-ones," <u>J. Chem. Soc. Perkin II</u> : 469-473 (1973)	
RA	A91	Coda et al., "(Z)- and (E)-Arylidene-1,3-dihydroindol-2-ones: Configuration, Conformation and Infrared Carbonyl Stretching Frequencies," <u>J. Chem. Soc. Perkin Trans. II</u> : 615-619 (1984)	
RA	A92	Coda et al., "3-(4-methylbenzylidene)-1,3-dihydroindol-2-one," <u>Journal of the Chemical Society, Perkin Transactions 2</u> 4:615-620 (1984) DATABASE CROSSFIRE, Beilstein Reference No. 6-21	
RA	A93	Decodts et al., "Suicide inhibitors of proteases. Lack of activity of halomethyl derivatives of some aromatic lactams," <u>Eur. J. Med. Chem</u> 18: 107-111 (1983)	
RA	A94	Desimoni et al., "Catalysis with Inorganic Cations. V ¹ Intramolecular Hetero Diels-Alder <i>versus</i> Ene Reactions: Effect of Magnesium perchlorate on Chemoselectivity," <u>Tetrahedron</u> 52(36) 12009-12018 (1196) © Pergamon	
RA	A95	Elliott and Rivers, "Reduction of Some Oxindolylidene Derivatives to 3-Substituted Oxindoles by Sodium Borohydride," <u>J. Med. Chem.</u> 29:2438-2440 (1964)	
RA	A96	Elliott et al., "1-methyl-2-(3-oxindolidenmethyl)-pyridinium," <u>Journal of Organic Chemistry</u> 29:2438-2440 (1964) DATABASE CROSSFIRE, Beilstein Reference No. 5-24	
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RA	A98	Hirao et al., "Rhodium-Catalyzed Carbonylation of 2-Alkynylaniline: Syntheses of 1,3-Dihydroindol-2-ones," <u>Tetrahedron Letters</u> 36(35) 1995 ©Pergamon	
RA	A99	Hodges et al., "Chemical and biological properties of some oxindolidyl-3-methines," <u>Canadian J. Chemistry</u> 46:2189-2194 (1968)	
*	A100	Howard, Harry R., "Lactam Derivatives," U.S. Provisional Patent Application Number 60/015134	
RA	A101	Howard et al., "Synthesis and aldose reductase inhibitory activity of substituted 2(1 <i>H</i>)-benzimidazolone- and oxindole-1-acetic acids," <u>Eur. J. Med. Chem.</u> 27:779-789 (1992) © Elsevier, Paris	
RA	A102	Katritzky et al., "Color and Constitution. Part 8[1]. Some Novel Dyestuffs Containing Indoxyl Residues," <u>J. Heterocyclic Chem.</u> 25:1287-1292 (1988)	
RA	A103	Kobayashi et al., "Anti-tumor Activity of Indole Derivatives," <u>Yakugaku Zasshi</u> 97:1033-1039 (1977)	
RA	A104	Kovac and Stetinova, "Furan derivatives. LXXX. Synthesis and properties of substituted furfurylidenoxindoles," <u>Chem. resu</u> 30:484-492 (1976)	
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LA	A106	Mariani et al., "Inhibition of angiogenesis by FCE 26806, a potent tyrosine kinase inhibitor," <u>Experimental Therapeutics - Proceedings of the American Association for Cancer Research</u> 35:381 at abstract no. 2268 (March 1994)	
DA	A107	Mohammadi et al., "Structures of the Tyrosine Kinase Domain of Fibroblast Growth Factor Receptor in Complex with Inhibitors," <u>Science</u> 276:955-960 (1997) © American Association for the Advancement of Science	
DA	A108	Neber and Röcker, "On the action of benzaldehydes on the free o-aminophenylacetic acid (II)," <u>Chem. Ber.</u> 56:1710-1716 (1923) (GERMAN AND ENGLISH TRANSLATION)	
DA	A109	Nodiff et al., "Antimalarial Phenanthrene Amino Alcohols. 3. Halogen-containing 9-phenanthrenemethanols," <u>Chemical Abstracts</u> , Vol. 83, abstract no. 188214 (1975)	
DA	A110	O'Sullivan and Rothery, "The Preparation and Possible Clinical Significance of 4'-Dialkylaminoisoidogenides," <u>Clinica Chimica Acta</u> 62:181-182 (1975) ©Elsevier Scientific Publishing Company	
DA	A111	Pavlenko et al., "Introduction of aminomethyl groups into heterocyclic CH-acid molecules," <u>Dopov. Akad. Nauk Ukr Rrsr, Ser. B: Geol., Khim. Biol. Nauki</u> 7:64-66 (1980)	
DA	A112	Plowman et al., "Receptor Tyrosine Kinases as Targets for Drug Intervention," <u>DN&P</u> 7:334-339 (1994)	
DA	A113	Quallich et al., "A General Oxindole Synthesis," <u>J. Synthetic Organic Chemistry</u> : 51-51 (1993)	
DA	A114	Schuchter et al., "Successful Treatment of Murine Melanoma with Bryostatin 1," <u>Cancer Research</u> 51:682-687 (1991)	
DA	A115	Shiraishi et al., "Specific inhibitors of Tyrosine-Specific Protein Kinase, Synthetic 4-Hydroxycinnamamide Derivatives," <u>Biochemical and Biophysical Research Communications</u> 147:322-328 (1987) © Academic Press	
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DA	A117	Singh et al., "Indolinone Derivatives as Potential Antimicrobial Agents," <u>Zentralbl. Mikrobiol.</u> 144:105-109 (1989) copyright VEB Gustav Fischer Verlag Jena	
DA	A118	Spada, et al., "Small molecule inhibitors of tyrosine kinase activity," <u>Expert Opinion on Therapeutic Patents</u> 5:805-817 (1995) ©Ashley Publications	
DA	A119	Sun et al., "Design, Synthesis, and Evaluations of Substituted 3-[(3- or 4-Carboxyethylpyrrol-2-yl)methylidene]indolin-2-ones as Inhibitors of VEGF, FGF, and PDGF Receptor Tyrosine Kinases," <u>Journal of Medicinal Chemistry</u> 42: 5120-5130 (1999) ©American Chemical Society	
DA	A120	Sun et al, "Synthesis and Biological Evaluations of 3-Substituted Indolin-2-ones: A Novel Class of Tyrosine Kinase Inhibitors That Exhibit Selectivity toward Particular Receptor Tyrosine Kinases," <u>J. Med. Chem.</u> 41:2588-2603 (1998) ©The American Chemical Society	
DA	A121	Tacconi and Marinone, "Preparazione e caratteristiche di alcuni 3-ossindolidenderivati," <u>Ricerca Scientifica</u> 38:1239-1244 (1968)	


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RA	A122	Tacconi et al., "(Z)- and (E)-3-Alkylidene-1,3-dihydroindol-2-ones: Influence of Configuration on the Transmission of the Inductive Effect to the Carbonyl Group," <u>J.C.S. Perkin II</u> 150-154 (1976)	
RA	A123	Thompson et al., "Facile Dimerisation of 3-Benzylideneindoline-2-thiones," <u>J. Chem. Soc. Perkin Trans. (I)</u> 1835-1837 (1993)	
RA	A124	Traxler, "Protein tyrosine kinase inhibitors in cancer treatment," <u>Expert Opinion on Therapeutic Patents</u> 7(6):571-588 (1997) © Ashley Publications Ltd.	
RA	A125	Wahl et al., "3-benzilidene-5-methyl-1,3-dihydroindol-2-one," <u>Ann. Chim.</u> 350 (1926), DATABASE CROSSFIRE, Beilstein Reference No. 2-21-00-00290	
RA	A126	Wahl, Beilstein Reg. No. 191439, <u>Bull. Soc. Chim. Fr.</u> , page 1038 (1909)	
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RA	A128	Walker et al., "Synthesis of New 3-(Pyridylmethylene)-, 3-(Pyridylmethyl)-, 3-(Piperidylmethyl)-, and 3-(β -Alkylaminoethyl)-2-indolinones. The Reduction of Isoindogenides, Nitro Compounds, and Pyridines in a Series of 2-Indolinones," <u>J. Med. Chem.</u> 8:626-637 (1965)	
RA	A129	Wright et al., "Cyclic Hydroxamic Acids Derived from Indole," <u>J. Am. Chem. Soc.</u> 78:221-224 (1956)	
RA	A130	Wright et al., "Inhibition of Angiogenesis in Vitro and In Ovo With an Inhibitor of Cellular Protein Kinases, MDL 27032," <u>J. Cellular Physiology</u> 152:448-457 (1992)	
RA	A131	Zhang et al., "Microtubule Effects of Welwistatin, a Cyanobacterial Indolinone that Circumvents Multiple Drug Resistance," <u>Molecular Pharmacology</u> 49:228-234 (1996) copyright The American Society for Pharmacology and Experimental Pharmaceutics	

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<i>RA</i>	A132	Zhungietu et al., "Reaction of Indoles and 2-Ketoindolines With Some Aldehydes," <u>Chemical Abstracts</u> , Vol. 78, abstract no. 111201 (1990)
<i>*</i>	A133	English Translation of Hungarian Patent No. 3899/92
	A134	English Translation of German Patent No. 2159360 (Ref. No. A85)
	A135	English Translation of German Patent No. 2159361 (Ref. No. A86)
	A136	English Translation of German Patent No. 2159363 (Ref. No. A88)
	A137	English Translation of German Patent No. 2321656 (Ref. No. A89)
	A138	English Translation of German Patent No. 3426419 (Ref. No. A90)
	A139	English Translation of European Patent No. 580502 (Ref. No. A96)
	A140	English Translation of European Patent No. 632102 (Ref. No. A98)
	A141	English Translation of French Patent No. 2689397 (Ref. No. A107)
	A142	English Translation of Japanese Patent No. 6229570 (Ref. No. A110)
	A143	English Translation of Japanese Patent No. 6239564 (Ref. No. A111)
	A144	English Translation of Japanese Patent No. 63141955 (Ref. No. A 112)
	A145	English Translation of Japanese Patent No. 558894 (Ref. No. A113)
	A146	English Translation of German Patent No. 878539 (Ref. No. A84)
	A147	English Translation of French Patent No. 1398224 (Ref. No. A105)
	A148	English Translation of French Patent No. 1599772 (Ref. No. A106)

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